#### **AMENDMENT TO THE CLAIMS**

# Claim 1 (currently amended)

A process for the preparation of <u>a salt of</u> an ω-benzyl ester of an amino dicarboxylic acid, comprising <u>A</u>) reacting the amino dicarboxylic acid with a benzyl alcohol derivative of the formula

$$(R^1)_n$$
  $CH_2OH$ 

wherein R<sup>1</sup> is hydrogen or is individually selected from the group consisting of hydrogen, C<sub>1</sub> to C<sub>4</sub> alkyl, C<sub>1</sub> to C<sub>4</sub> alkoxy and halogen and n is 1 or 3, in the presence of an alkanesulfonic acid catalyst with a molar ratio of at least one mole of catalyst per mole of the amino dicarboxylic acid of an alkanesulfonic acid, B) optionally in the presence of a solvent to obtain a salt thereof and optionally reacting the salt with a base to obtain the free form thereof.

# Claim 2 (previously presented)

The process of Clam 1, wherein the amino dicarboxylic acid is an  $\alpha$ -amino carboxylic acid carrying another carboxyl group attached to a carbon other than that in the  $\alpha$  position.

## Claim 3 (previously presented)

The process of Claim 2, wherein the amino dicarboxylic acid is glutamic acid or aspartic acid.

# Claim 4 (previously presented)

The process of claim 1 wherein the alcohol of formula (I) is benzyl alcohol.

#### Claim 5 (currently amended)

The process of claim 1 wherein the temperature of the  $\underline{A}$ ) reaction is less than or equal to 80°C.

#### Claim 6 (previously presented)

The process of claim 1 wherein the benzyl alcohol or its derivative of formula (I) is used in an amount of 1.2 to 3 mol per mole of the amino dicarboxylic acid.

#### Claim 7 (previously presented)

The process of claim 1 wherein the alkanesulfonic acid is methanesulfonic acid.

## Claim 8 (previously presented)

The process of claim 1 wherein the amount of alkanesulfonic acid used is 1.01 to 2 mol per mole of the amino dicarboxylic acid.

## Claim 9 (currently amended)

The process of claim 1 wherein the solvent of the <u>A)</u> reaction is selected from the group consisting of aliphatic and aromatic and halogenated and nonhalogenated hydrocarbons.

#### Claim 10 (previously presented)

The process of claim 1 wherein the  $\omega$ -benzyl ester of the amino dicarboxylic acid is obtained in the free form by bringing the alkanesulfonate of the  $\omega$ -benzyl ester of the amino dicarboxylic acid into contact with an organic or inorganic base.

# Claim 11 (previously presented)

The process of Claim 10, wherein the base is used in an amount sufficient to reach the isoelectric point of the ester to be obtained.

#### Claim 12 (previously presented)

The process of Claim 10 wherein the base is an aqueous ammonia solution.

## Claim 13 (currently amended)

The process of claim 1 wherein the <u>salt</u> alkanesulfonate of the  $\omega$ -benzyl ester of the amino dicarboxylic acid is crystallized before being converted to the free  $\omega$ -benzyl ester of the amino dicarboxylic acid.

## Claim 14 (currently amended)

The process of claim 1 wherein a solvent/water azeotrope is distilled off the A) reaction at a temperature of less than 80°C.

# Claim 15 (currently amended)

The process of claim 1 wherein the <u>salt</u> alkanesulfonate of the ω benzyl ester of the amino dicarboxylic acid is isolated before being brought into contact with the base.

#### Claim 16 (currently amended)

The process of claim 1 wherein the alkanesulfonate of the  $\omega$  benzyl-ester of the amino dicarboxylic acid salt is not isolated from the medium before this ester is released.

#### Claim 17 (currently amended)

The process of claim 1 wherein the <u>salt alkanesulfonate of the ω benzyl ester to</u> be converted to the free ester is dissolved in water.

# Claim 18 (previously presented)

The process of claim 17 wherein a solvent for the benzyl alcohol derivative is added with the water or after the introduction of water.

## Claim 19 (currently amended)

The process of claim 1 wherein, after having reached the pH of the isoelectric point of the A) reaction, the medium is heated.

Claim 20 (cancelled)

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Claim 21 (previously presented)

Alkanesulfonate having the formula

wherein  $R^1$  is hydrogen or is individually selected from the group consisting of  $C_1$  to  $C_4$  alkyl,  $C_1$  to  $C_4$  alkoxy and halogen, A is selected from the group consisting of an aliphatic, cycloaliphatic, aryl, araliphatic or heterocyclic and  $R^2$  is alkane residue of the alkanesulfonic acid.

# Claim 22 (previously presented)

An alkanefulsonate of claim 21 wherein it is  $\gamma$ -benzyl glutamate methanesulfonate or  $\beta$ -benzyl aspirate methane sulfonate.